



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICANT : FARRELL, David H.
SERIAL NO : 10/006,403
FILED : December 5, 2001
TITLE : A MONOCLONAL ANTIBODY-BASED DIAGNOSTIC ASSAY FOR
GAMMA FIBRINOGEN

Grp./A.U. : 1632
Conf. No. : 4264
Docket No. : P05375US1

RESPONSE TO OFFICE COMMUNICATION

Assistant Commissioner for Patents
Washington, D.C. 20231

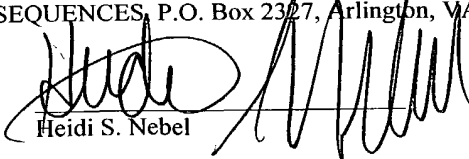
Dear Sir:

In response to the office communication dated April 8, 2002, Paper No. 12, Applicant submits herewith is a duplicate copy of the sequence listing earlier submitted March 4, 2002.

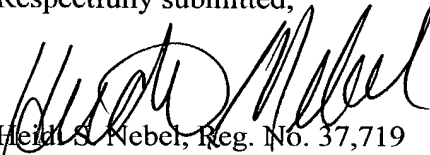
No fees or extensions of time are believed to be due in connection with this amendment; however, consider this a request for any extension inadvertently omitted, and charge any additional fees to Deposit Account No. 26-0084.

CERTIFICATE OF MAILING

I hereby certify that this correspondence is being mailed via first class mail, in an envelope addressed to: United States Patent and Trademark Office, BOX SEQUENCES, P.O. Box 2327, Arlington, VA 22202, this 15 day of May, 2002.


Heidi S. Nebel

Respectfully submitted,



Heidi S. Nebel, Reg. No. 37,719

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APPLICATION NUMBER	FILING/RECEIPT DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NUMBER
10/006,403	12/05/2001	David H. Farrell	P05375US1

27407
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CONFIRMATION NO. 4264

FORMALITIES LETTER



OC000000007824117

Date Mailed: 04/08/2002

**NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS
CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE
DISCLOSURES**


Applicant is given **TWO MONTHS FROM THE DATE OF THIS NOTICE** within which to file the items indicated below to avoid abandonment. Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

- The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d). Applicant must provide a substitute computer readable form (CRF) copy of the "Sequence Listing" and a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter, as required by 37 CFR 1.821(e), 1.821(f), 1.821(g), 1.825(b), or 1.825(d). If applicant desires the sequence listing in the instant application to be identical with that of another application on file in the U.S. Patent and Trademark Office, such request in accordance with 37 CFR 1.821(e) may be submitted in lieu of a new CRF.

For questions regarding compliance to these requirements, please contact:

- For Rules Interpretation, call (703) 308-4216
- To Purchase PatentIn Software, call (703) 306-2600
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*A copy of this notice **MUST** be returned with the reply.*


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PART 2 - COPY TO BE RETURNED WITH RESPONSE